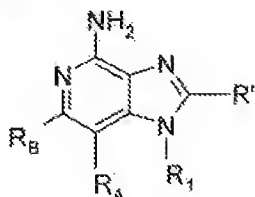


Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (currently amended) A compound of the following Formula I:



I

wherein:

R₁ has the formula alkylene-L-R₁₋₁, alkenylene-L-R₁₋₁, or alkynylene-L-R₁₋₁, wherein:

the alkylene, alkenylene, and alkynylene groups are optionally interrupted with one or more -O- groups;

L is a bond or a functional linking group selected from the group consisting of -NH-S(O)₂-, -NH-C(O)-, -NH-C(S)-, -NH-S(O)₂-NR₃-, -NH-C(O)-NR₃-, -NH-C(S)-NR₃-, -NH-C(O)-O-, -O-, -S-, and -S(O)₂-; and

R₁₋₁ is a linear or branched aliphatic group having at least 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;

R'' is selected from the group consisting of:

_____ hydrogen;

_____ alkyl;

_____ alkenyl;

_____ aryl;

_____ heteroaryl;

_____ heterocyclyl;

_____ alkylene-Y-alkyl;

_____ alkylene-Y-alkenyl;

alkylene-Y-aryl; and

alkyl or alkenyl substituted by one or more substituents selected from the group consisting of:

_____ -OH;
 _____ halogen;
 _____ -N(R₄)₂;
 _____ -C(O)-C₁₋₁₀alkyl;
 _____ -C(O)-O-C₁₋₁₀alkyl;
 _____ -N₃;
 _____ -aryl;
 _____ heteroaryl;
 _____ heterocyclyl;
 _____ -C(O)-aryl; and
 _____ -C(O)-heteroaryl;

wherein: Y is -O- or -S(O)₀₋₂; and each R₄ is independently selected from the group consisting of hydrogen, C₁₋₁₀alkyl, and C₂₋₁₀alkenylhydrogen or a non-interfering substituent;

R_A and R_B are each independently selected from the group consisting of:

hydrogen,
 halogen,
 alkyl,
 alkenyl,
 alkoxy,
 alkylthio, and
 -N(R₃)₂;

or when taken together, R_A and R_B form a fused aryl ring or heteroaryl ring containing one heteroatom selected from the group consisting of N and S wherein the aryl or heteroaryl ring is unsubstituted or substituted by one or more R groups;

or when taken together, R_A and R_B form a fused 5- to 7-membered saturated ring, optionally containing one heteroatom selected from the group consisting of N and S, and unsubstituted or substituted by one or more R groups;

each R is independently selected from the group consisting of

_____halogen,

_____hydroxy,

_____alkyl,

_____alkenyl,

_____haloalkyl,

_____alkoxy,

_____alkylthio, and

_____N(R₃)₂R_A and R_B form a fused aryl ring or heteroaryl ring containing one heteroatom or a fused 5- to 7-membered saturated ring, optionally containing one heteroatom, wherein the heteroatom is selected from the group consisting of N and S, and wherein the aryl, heteroaryl, or 5- to 7-membered saturated ring is unsubstituted or substituted by one or more non-interfering substituents; and

each R₃ is independently selected from the group consisting of hydrogen and alkyl;

with the proviso that when L is -NH-S(O)₂- and R_A and R_B join to form an unsubstituted benzene ring, R₁₋₁ is a linear or branched aliphatic group having greater than 16 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds; and with the further proviso that when L is -NH-C(O)- and R_A and R_B join to form an unsubstituted pyridine ring, R₁₋₁ is a linear or branched aliphatic group having greater than 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;
or a pharmaceutically acceptable salt thereof.

2-6 (canceled)

7. (currently amended) The compound or salt of claim 16 wherein R_A and R_B form a fused benzene ring which is unsubstituted.

8-10 (canceled)

11. (currently amended) The compound or salt of claim 10 wherein L is a bond or a functional linking group selected from the group consisting of -NH-C(O)-, -NH-S(O)₂-, and -NH-C(O)-N(R₃)₂-.

12 (canceled)

13. (currently amended) The compound or salt of claim 12 wherein R_{1-1} is a linear or branched aliphatic group having 12-20 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds.

14. (original) The compound or salt of claim 13 wherein R_{1-1} is a straight chain C_{12} - C_{20} alkyl.

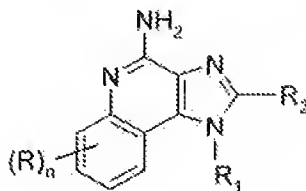
15-16 (canceled)

17. (currently amended) The compound or salt of claim 16 wherein R_1 has the formula C_{1-5} alkylene-L- R_{1-1} and the C_{1-5} alkylene is optionally interrupted with one -O- group.

18. (currently amended) The compound or salt of claim 15 wherein R_2 is selected from the group consisting of hydrogen, alkyl, and alkylene-O-alkyl.

19 (canceled)

20. (original) A compound of the following Formula III:



III.

wherein:

R_1 has the formula alkylene-L- R_{1-1} , alkenylene-L- R_{1-1} , or alkynylene-L- R_{1-1} , wherein:

the alkylene, alkenylene, and alkynylene groups are optionally interrupted with one or more -O- groups;

L is a bond or a functional linking group selected from the group consisting of $\text{-NH-S(O)}_2\text{-}$, -NH-C(O)- , -NH-C(S)- , $\text{-NH-S(O)}_2\text{-NR}_3\text{-}$, $\text{-NH-C(O)-NR}_3\text{-}$, $\text{-NH-C(S)-NR}_3\text{-}$, -NH-C(O)-O- , -O- , -S- , and $\text{-S(O)}_2\text{-}$; and

R_{1-4} is a linear or branched aliphatic group having at least 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;

R is selected from the group consisting of

halogen,
hydroxy,
alkyl,
alkenyl,
haloalkyl,
alkoxy,
alkylthio, and
 $\text{-N(R}_3)_2$;

n is 0 to 4;

R_2 is selected from the group consisting of:

hydrogen;
alkyl;
alkenyl;
aryl;
heteroaryl;
heterocyclyl;
alkylene-Y-alkyl;
alkylene-Y-alkenyl;
alkylene-Y-aryl; and

alkyl or alkenyl substituted by one or more substituents selected from the group consisting of:

-OH ;
halogen;
 $\text{-N(R}_4)_2$;
 $\text{-C(O)-C}_{1-10}\text{alkyl}$;

-C(O)-O-C₁₋₁₀alkyl;

-N₃;

aryl;

heteroaryl;

heterocyclyl;

-C(O)-aryl; and

-C(O)-heteroaryl;

Y is -O- or -S(O)₀₋₂;

each R₄ is independently selected from the group consisting of hydrogen, C₁₋₁₀alkyl, and C₂₋₁₀alkenyl; and

R₃ is selected from the group consisting of hydrogen and alkyl;

with the proviso that when L is -NH-S(O₂)-, and n is 0, R₁₋₁ is a linear or branched aliphatic group having at least 16 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;
or a pharmaceutically acceptable salt thereof.

21. (original) The compound or salt of claim 20 wherein n is 0.

22-23 (canceled)

24. (currently amended) A pharmaceutical composition comprising a therapeutically effective amount of a compound or salt of ~~any one of claims 1 through 23~~ in combination with a pharmaceutically acceptable carrier.

25. (currently amended) A method of inducing cytokine biosynthesis in an animal comprising administering an effective amount of a compound or salt of ~~any one of claims 1 through 23~~ to the animal.

26-27 (canceled)

28. (currently amended) A method of vaccinating an animal comprising administering an effective amount of a compound or salt of ~~any one of claims 1 through 23~~ to the animal as a vaccine adjuvant.

29. (currently amended) A method of vaccinating an animal comprising administering an effective amount of *N*-(2-{2-[4-amino-2-(2-methoxyethyl)-1*H*-imidazo[4,5-*c*]quinolin-1-yl]ethoxy}ethyl)hexadecanamide to the animal as a vaccine adjuvant.

30-32 (canceled)

33. (new) A pharmaceutical composition comprising a therapeutically effective amount of a compound or salt of claim 20 in combination with a pharmaceutically acceptable carrier.

34. (new) A method of inducing cytokine biosynthesis in an animal comprising administering an effective amount of a compound or salt of claim 20 to the animal.

35. (new) A method of vaccinating an animal comprising administering an effective amount of a compound or salt of claim 20 to the animal as a vaccine adjuvant.